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DATE MAILED: 09/30/2003

APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
09/763,994	02/27/2001	Edmonds Taylor Brian	X-12239	6826
75	590 09/30/2003			
Robert L Sharp Eli Lilly & Company Lilly Corporate Center DC1104			EXAMINER	
			ROMEO, DAVID S	
Indianapolis, IN	N 46285		ART UNIT	PAPER NUMBER

Please find below and/or attached an Office communication concerning this application or proceeding.

		Application No.	Applicant(s)			
		09/763,994	BRIAN, EDMONDS TAYLOR			
Office Action Summary		Examiner	Art Unit			
		DAVID ROMEO	1647			
Period fo	- The MAILING DATE of this communication app r Reply	oears on the cover sheet with the	e correspondence address			
THE M - Exten after: - If the - If NO - Failur - Any re	DRTENED STATUTORY PERIOD FOR REPL (AAILING DATE OF THIS COMMUNICATION. solons of time may be available under the provisions of 37 CFR 1.1 SIX (6) MONTHS from the mailing date of this communication. Period for raply specified above is less than thirty (30 days, a raple period for raply specified above is loss than thirty (30 days, a raple to the communication of the communication	(36(a). In no event, however, may a reply be by within the statutory minimum of thirty (30) of will apply and will expire SIX (6) MONTHS fr s, cause the application to become ABANDO	timely filed days will be considered timely. on the mailing date of this communication. NED (35 U.S.C. § 133).			
1)⊠	Responsive to communication(s) filed on 27	February 2001 .				
2a)□	This action is FINAL . 2b)⊠ Th	nis action is non-final.				
3)[_ Dispositi	Since this application is in condition for allow closed in accordance with the practice under on of Claims	ance except for formal matters, Ex parte Quayle, 1935 C.D. 11	prosecution as to the merits is , 453 O.G. 213.			
4)⊠	Claim(s) 32-47 is/are pending in the application	on.				
4a) Of the above claim(s) is/are withdrawn from consideration.						
5) Claim(s) is/are allowed.						
6) Claim(s) is/are rejected.						
7)	7) Claim(s) is/are objected to.					
8)🖂	Claim(s) 32-47 are subject to restriction and/o	r election requirement.				
Applicati	on Papers					
9)☐ The specification is objected to by the Examiner.						
10)☐ The drawing(s) filed on is/are: a)☐ accepted or b)☐ objected to by the Examiner.						
Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).						
11) The proposed drawing correction filed on is: a) approved b) disapproved by the Examiner.						
If approved, corrected drawings are required in reply to this Office action.						
12)☐ The oath or declaration is objected to by the Examiner.						
Priority u	nder 35 U.S.C. §§ 119 and 120					
13) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).						
a)[☐ All b)☐ Some * c)☐ None of:					
	1. Certified copies of the priority document	ts have been received.				
	2. Certified copies of the priority documents have been received in Application No					
	Copies of the certified copies of the prior application from the International Bure the attached detailed Office action for a list.	reau (PCT Rule 17.2(a)).	_			
	cknowledgment is made of a claim for domest	·				
_a)	☐ The translation of the foreign language procknowledgment is made of a claim for domest	ovisional application has been r	eceived.			
Attachment		,				
2) Notice	o of References Cited (PTO-892) o of Draftsperson's Patent Drawing Review (PTO-948) nation Disclosure Statement(s) (PTO-1449) Paper No(s)	5) Notice of Inform	ary (PTO-413) Paper No(s) al Patent Application (PTO-152)			

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DETAILED ACTION

Election/Restrictions

- Restriction is required under 35 U.S.C. 121 and 372.
- This application contains the following inventions or groups of inventions which are not so linked as to form a single general inventive concept under PCT Rule 13.1.
- In accordance with 37 CFR 1.499, applicant is required, in reply to this action, to elect a single invention to which the claims must be restricted.

Group 1, claim(s) 32-36 and 38, drawn to an <a href="https://h

Group 2, claim(s) 37 and 39, drawn to an antibody and compositions comprising same.

Group 3, claim(s) 40 and 43, drawn to a method for inhibiting or stimulating tissue growth which comprises administering to a patient in need thereof a polypeptide.

Group 4, claim(s) 41 and 42, drawn to a method for inhibiting tissue growth which comprises administering to a patient in need thereof an *antibody*.

Group 5, claim(s) 44, drawn to a method for modulating a <u>TGFβ regulatable activity</u> comprising administering an *antibody*.

Group 6, claim(s) 45, drawn to a method for modulating a <u>TGFβ regulatable activity</u> comprising administering an *antisense nucleic acid*.

Group 7, claim(s) 46, drawn to a method for prevention and/or treatment comprising administering a *polypeptide*.

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Group 8, claim(s) 47, drawn to a method for prevention and/or treatment comprising administering an *antibody*.

4. The inventions listed as Groups 1-9 do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding special technical features for the following reasons:

Group 1 recites the special technical feature of a nucleic acid encoding a polypeptide, which is not required by any of the other Groups.

Group 2 recites the special technical feature of an antibody, which is not required by any of the other Groups.

Group 3 recites the special technical feature of stimulating tissue growth, which is not required by any of the other Groups.

Group 4 recites the special technical feature of inhibiting tissue growth by administering an antibody, which is not required by any of the other Groups.

Group 5 recites the special technical feature of modulating TGF β regulatable using an antibody, which is not required by any of the other Groups.

Group 6 recites the special technical feature of an antisense nucleic acid, which is not required by any of the other Groups.

Group 7 recites the special technical feature of using a polypeptide as a therapeutic agent, which is not required by any of the other Groups.

Group 8 recites the special technical feature of using an antibody as a therapeutic agent, which is not required by any of the other Groups.

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5. This application contains claims directed to more than one species of the generic invention. These species are deemed to lack unity of invention because they are not so linked as to form a single general inventive concept under PCT Rule 13.1.

- 6. In order for more than one species to be examined, the appropriate additional examination fees must be paid. The species are as follows:
 - a. SEQ ID NO: 2
 - b. SEQ ID NO: 4
 - c. SEQ ID NO: 5
 - d. SEQ ID NO: 6
- If applicant selects any one of Groups 1-8, one species from the SEQ ID NO must be chosen to be fully responsive.
- 8. In order for more than one species to be examined, the appropriate additional examination fees must be paid. The species are as follows:
 - e. Cancer
 - f. Fibrosis
 - g. Osteoporosis
 - h. Myocardial infarction
 - Congestive heart failure
 - j. Dilated cardiomyopathy
 - k. Deep venous thrombosin
 - Disseminated intravascular thrombosis
 - m. Stroke

n. Sepsis

- o. Injuries involving major tissue damage and trauma
- p. Systemic inflammatory response syndrome
- q. Sepsis syndrome
- r. Septic shock
- Multiple organ dysfunction syndrome (including DIC)
- t. Atherosclerotic plaque rupture
- u. Associated sequela
- If applicant selects either Groups 7 or 8, one species from the disease group must be chosen to be fully responsive.
- Applicant is advised that the reply to this requirement to be complete must include an
 election of the invention to be examined even though the requirement be traversed (37 CFR
 1.143).
- 11. Because these inventions are distinct for the reasons given above and have acquired a separate status in the art because of their recognized divergent subject matter, separate search requirements, and/or different classification, restriction for examination purposes as indicated is proper.
- 12. Applicant is reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 CFR 1.48(b) if one or more of the currently named inventors is no longer an inventor of at least one claim remaining in the

application. Any amendment of inventorship must be accompanied by a request under 37 CFR 1.48(b) and by the fee required under 37 CFR 1.17(i).

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Conclusion

Any inquiry concerning this communication or earlier communications from the examiner should be directed to **David Romeo**, **Ph.D**. whose telephone number is 703-305-4050. The examiner can normally be reached on Monday through Friday, 8:00AM to 5:00PM.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, **Gary Kunz, Ph.D.** can be reached on 703-308-4623. The fax phone numbers for the organization where this application or proceeding is assigned are 703-872-9306 for regular communications and 703-872-9307 for After Final communications. The fax phone numbers for the customer service center is 703-872-9305

Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to the receptionist whose telephone number is 703-308-0196.

CJN September 29, 2003 SUPERVISORY PATENT EXAMINER
TECHNOLOGY CENTER 1800